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Abstract of the Disclosure

The dihydroindole C-ring found in CC-1065/duocarmycin analogs is formed by the 5-*exo-trig* radical cyclization of an aryl halide onto a tethered vinyl chloride forming with chlorine installed as a suitable leaving group for subsequent cyclopropane spirocyclization. The versatility of this approach is disclosed in the context of six CC-1065 / duocarmycin analogs previously synthesized in this laboratory.

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